

AMENDMENTS TO THE CLAIMS

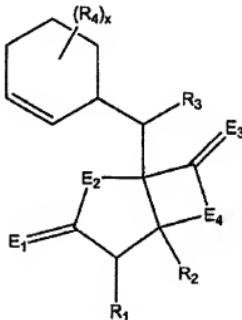
Please amend claims 1, 4, 9-10, 13-21, and 23-24 as indicated below.

Please cancel claim 11 without prejudice.

The listing of claims will replace all prior versions, and listings, of claims in the application.

Listing of Claims:

1. (Currently Amended) An isolated compound having of the structure (I):



(I)

wherein:

R_1 is a substituted alkyl;

R_2 is methyl;

R_3 is hydroxy;

Each R₄ is independently alkyl, substituted alkyl, alkenyl, substituted alkenyl, alkynyl, substituted alkynyl, aryl, substituted aryl, cycloalkyl, or substituted cycloalkyl;

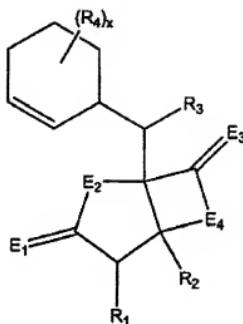
E₁, E₃, and E₄ are—O;

E₂ is —NR₅, wherein R₅ is —H or C₁-C₆ alkyl; and

X is 0 to 8.

2. (Previously Presented) The compound of Claim 1, wherein E₂ is -NH.
3. (Canceled).
4. (Currently Amended) The compound of Claim 1, wherein R₁ is a substituted alkyl substituted with one or more substitutions substituents selected from the group consisting of halogen, cyano, oxyacyl, amino, amido, -C(O)H, acyl, carboxyl, and sulfonamide.
5. (Original) The compound of Claim 4, wherein the substituted alkyl is a halogenated alkyl.
6. (Original) The compound of Claim 5, wherein the halogenated alkyl is a chlorinated alkyl.
7. (Previously Presented) A pharmaceutical composition comprising at least one compound of Claim 1 in a pharmaceutically acceptable carrier thereof.
8. (Canceled).
9. (Currently Amended) The pharmaceutical composition of Claim 7, further comprising at least one additional anti-neoplastic agent in combination with the at least one compound of claim 1.

10. (Currently Amended) A method of treating a human mammalian cancer cell proliferative disorder, comprising administering to a subject in need thereof contacting human refractile cancer cells with a therapeutically effective amount of a compound having of the structure (I):



(I)

wherein:

R₁ to R₃ are each independently -H, alkyl, substituted alkyl, alkenyl, substituted alkenyl, alkynyl, substituted alkynyl, aryl, substituted aryl, heteroaryl, substituted heteroaryl, heterocyclic, substituted heterocyclic, cycloalkyl, substituted cycloalkyl, alkoxy, substituted alkoxy, thioalkyl, substituted thioalkyl, hydroxy, halogen, amino, amido, carboxyl, -C(O)H, acyl, oxyacyl, carbamate, sulfonamide, or sulfonyl;

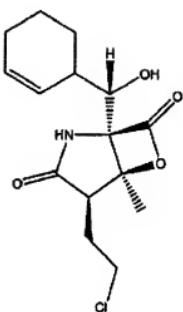
Each R₄ is independently alkyl, substituted alkyl, alkenyl, substituted alkenyl, alkynyl, substituted alkynyl, aryl, substituted aryl, cycloalkyl, or substituted cycloalkyl;

E₁ to E₄ are each independently -O-, NR₅, or -S, wherein R₅ is -H or C₁-C₆ alkyl; and

X is 0 to 8.

11. (Canceled).

12. (Currently Amended) The method of Claim 10, wherein the compound has is of the structure of formula (V):



(V).

13. (Currently Amended) The method of Claim 10, wherein the human refractile cancer cell proliferative disorder is selected from the group consisting of a mammary cancer cell, a small-cell lung cancer cell, a non-small-cell lung cancer cell, a colorectal cancer cell, a leukemia cancer cell, a melanoma cancer cell, a pancreatic adenocarcinoma cancer cell, a central nervous system (CNS) cancer cell, an ovarian cancer cell, a prostate cancer cell, a sarcoma cell of a soft tissue cancer, a sarcoma cell of a bone, a head cancer cell, a neck cancer cell, a gastric cancer cell, a thyroid cancer cell, a stomach cancer cell, a myeloma cancer cell, a bladder cancer cell, a renal cancer cell, a neuroendocrine cancer cell, a non-Hodgkin's cancer disease, cell and a Hodgkin's disease cancer cell.

14. (Currently Amended) The method of Claim 13, wherein the human refractile cancer cell proliferative disorder is a sarcoma cell of a soft tissue or a sarcoma cell of a bone.
15. (Currently Amended) The method of Claim 13, wherein the human refractile cancer cell proliferative disorder is a leukemia cancer cell.
16. (Currently Amended) The method of Claim 13, wherein the human refractile cancer cell proliferative disorder is a myeloma cancer cell.
17. (Currently Amended) The method of Claim 13, wherein the human refractile cancer cell proliferative disorder is an ovarian cancer cell.
18. (Currently Amended) The method of Claim 13, wherein the human refractile cancer cell proliferative disorder is a prostate cancer cell.
19. (Currently Amended) The method of Claim 13, wherein the human refractile cancer cell proliferative disorder is a non-Hodgkin's disease cancer cell.
20. (Currently Amended) The method of Claim 13, wherein the human refractile cancer cell proliferative disorder is a pancreatic adenocarcinoma cancer cell.
21. (Currently Amended) The method of Claim 10, wherein: R₁ is a substituted alkyl; R₂ is methyl; R₃ is hydroxy; each R₄ is independently alkyl, substituted alkyl, alkenyl, substituted alkenyl, alkynyl, substituted alkynyl, aryl, substituted aryl, cycloalkyl, or substituted cycloalkyl; E₁, E₄ and E₃ are -O; E₂ is -NR₅, wherein R₅ is -H or C₁-C₆ alkyl; and X is 0 to 8.
22. (Previously Presented) The method of Claim 21, wherein R₁ is a halogenated alkyl.
23. (Currently Amended) The method of Claim 10, further comprising administering at least one additional anti-neoplastic agent in combination with at least one compound of the structure (I).

In re Application of

William H. Fenical et al.

Application No.: 10/561,711

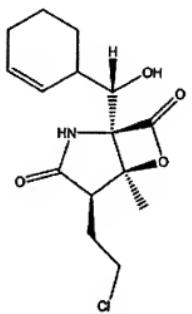
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24. (Currently Amended) A pharmaceutical composition comprising a pharmaceutically acceptable carrier and an effective amount of a compound having of the structure:



wherein the pharmaceutical composition is in a solid form.

25. (Previously Presented) The compound of Claim 1, wherein the structure is:

